AMENDMENTS TO THE CLAIMS

The following listing of claims will replace all prior versions and listings of claims in the application.

LISTING OF CLAIMS

What is claimed is:

- 1. (currently amended) An arteriovenous shunt comprising:
- a. an arterial graft comprising a body, a lead end and a terminal end, wherein said lead end being configured is operable for subcutaneous connection to an artery by anastomosis; and
- b. a single lumen venous outflow catheter comprising an intake end and depositing end, wherein said depositing end being configured is operable for insertion through a vein into the right atrium of the heart; and
- c. a cuff <u>operable to direct passage of blood from said arterial graft to said</u>

 <u>venous outflow catheter, said cuff</u> comprising an inlet <u>in fluid communication with</u> and an outlet, wherein:
 - i. said inlet being is connected to said terminal end of said arterial subcutaneous graft; and
 - ii. said outlet being is connected to said intake end of said venous outflow catheter.
- 2. (currently amended) The arteriovenous shunt of claim 1 wherein said <u>arterial</u> subcutaneous graft is made of a biocompatible flexible material.

- 3. (original) The arteriovenous shunt of claim 2, wherein said biocompatible flexible material is polytetrafluoroethylene (PTFE) or polyurethane.
- 4. (original) The arteriovenous shunt of claim 1, wherein said arterial graft has a diameter from about 2 mm to about 8 mm and a length from about 20 cm to about 60 cm.
- 5. (original) The arteriovenous shunt of claim 4, wherein said arterial graft has a diameter of from about 6 mm to about 8 mm and a length of about 40 cm.
- 6. (original) The arteriovenous shunt of claim 1, wherein said artery is the brachial, axillary, femoral or external iliac artery.
- 7. (currently amended) The arteriovenous shunt of claim 1, wherein said cuff is polytetrafluoroethylene Teflon or polyethylene terephthalate Dacron.
- 8. (original) The arteriovenous shunt of claim 1, wherein said venous outflow catheter has a diameter from about 1 mm to about 7 mm and a length of from about 20 cm to about 80 cm.
- 9. (original) The arteriovenous shunt of claim 1, wherein said venous outflow catheter has a diameter from about 5 mm to about 7 mm and a length of from about 40 cm to about 60 cm.

- 10. (original) The arteriovenous shunt of claim 1, wherein said venous outflow catheter is made of polyurethane or silicone.
- 11. (original) The arteriovenous shunt of claim 1, wherein said vein is the cephalic, axillary, jugular, femoral or external iliac vein.
- 12. (currently amended) The arteriovenous shunt of claim 1, wherein said venous outflow catheter has a diameter of about 1 mm smaller than said arterial the subcutaneous graft.
- 13. (currently amended) A system for performing hemodialysis on a patient comprising:
 - a. an arteriovenous shunt comprising:
 - i. an arterial graft comprising a body, a lead end and a terminal end,

 wherein said lead end being configured is operable for subcutaneous

 connection to an artery by anastomosis; and
 - ii. a single lumen venous outflow catheter comprising an intake end and depositing end, wherein said depositing end being configured is operable for insertion through a vein into the right atrium of the heart; and
 - iii. ii. a cuff operable to direct passage of blood from said arterial graft to said venous outflow catheter, said cuff comprising an inlet in fluid communication with and an outlet, wherein:

- said inlet <u>being</u> is connected to said terminal end of said subcutaneous graft; and
- said outlet <u>being</u> is connected to said intake end of said venous outflow catheter;

and

- b. a hemodialysis apparatus.
- 14. (currently amended) The system according to claim 13, wherein said venous outflow catheter has a diameter of about 1 mm smaller than said <u>arterial</u> subcutaneous graft.
- 15. (original) The system according to claim 13, wherein said artery is the brachial, axillary, femoral or external iliac artery.
- 16. (original) The system according to claim 13, wherein said vein is the cephalic, axillary, jugular, femoral or external iliac vein.
- 17. (currently amended) A method of performing hemodialysis on a patient comprising:
- a. <u>surgically</u> inserting an arteriovenous shunt into a patient, wherein said arteriovenous shunt comprises:
 - i. an arterial graft comprising a body, a lead end and a terminal end,

 wherein said lead end being configured is operable for subcutaneous

 connection to an artery by anastomosis; and

- ii. a single lumen venous outflow catheter comprising an intake end and depositing end, wherein said depositing end being configured is operable for insertion through a vein into the right atrium of the heart; and
- iii. a cuff operable to direct passage of blood from said arterial graft to said venous outflow catheter, said cuff comprising an inlet in fluid communication with and an outlet, wherein:
 - said inlet <u>being</u> is connected to said terminal end of said <u>arterial</u>
 subcutaneous graft; and
 - said outlet <u>being</u> is connected to said intake end of said venous outflow catheter;
- b. connecting said arterial graft to a hemodialysis apparatus;
- c. collecting blood from the patient through said arterial subcutaneous graft;
- d. passing said blood through the hemodialysis apparatus;
- e. collecting purified blood from hemodialysis apparatus; and
- f. transmitting said purified blood through said cuff into said venous outflow catheter.
- 18. (currently amended) The method according to claim 16 wherein said venous outflow catheter has a diameter of about 1 mm smaller than said <u>arterial</u> subcutaneous graft.
- 19. (original) The method according to claim 16, wherein said artery is the brachial, axillary, or femoral, external iliac artery.

	20.	(original) The method according to clai	m 16, wherein said vein is the axillary
jugular, femoral or external iliac vein.			